

4. A method according to Claim 3, wherein said dedimethylaminotetracycline is selected from the group consisting of 4-dedimethylaminotetracycline, 4-dedimethylamino-5-oxytetracycline, 4-dedimethylamino-7-chlorotetracycline, 4-hydroxy-4-dedimethylaminotetracycline, 5a,6-anhydro-4-hydroxy-4-dedimethylaminotetracycline, 6  $\alpha$ -deoxy-5-hydroxy-4-dedimethylaminotetracycline, 6-demethyl-6-deoxy-4-dedimethylaminotetracycline, 4-dedimethylamino-12a-deoxytetracycline, 12  $\alpha$ -deoxy-4-deoxy-4-dedimethylaminotetracycline, 12a, 4  $\alpha$ -anhydro-4-dedimethylaminotetracycline, 7-dimethylamino-6-demethyl-6-deoxy-4-dedimethylaminotetracycline, 5-hydroxy-6-  $\alpha$ -deoxy-4-dedimethylaminotetracycline, 4-dedimethylamino-12  $\alpha$ -deoxyanhydrotetracycline and 4-dedimethylamino-11-hydroxy-12a-deoxytetracycline.

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$\beta^1$  10. (Amended) A method according to Claim 1, wherein said tetracycline derivative is doxycycline.

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11. A method according to Claim 1, wherein said tetracycline derivative is administered systemically.

12. A method according to Claim 11, wherein said tetracycline derivative is administered systemically by a controlled release delivery system.

13. A method according to Claim 1, wherein said tetracycline derivative is administered orally.

14. A method according to Claim 1, wherein said tetracycline derivative is administered topically.

18. A method of treating cataract formation in a mammal comprising administering to the mammal an effective amount of a tetracycline derivative.

19. A method according to Claim 18, wherein said tetracycline derivative is a non-antimicrobial tetracycline.

20. A method according to Claim 18, wherein said tetracycline derivative is a dedimethylaminotetracycline.

21. A method according to Claim 20, wherein said dedimethylaminotetracycline is selected from the group consisting of 4-dedimethylaminotetracycline, 4-dedimethylamino-5-oxytetracycline, 4-dedimethylamino-7-chlorotetracycline, 4-hydroxy-4-dedimethylaminotetracycline, 5a,6-anhydro-4-hydroxy-4-dedimethylaminotetracycline, 6  $\alpha$  -deoxy-5-hydroxy-4-dedimethylaminotetracycline, 6-demethyl-6-deoxy-4-dedimethylaminotetracycline, 4-dedimethylamino-12a-deoxytetracycline, 12 $\alpha$  -deoxy-4-deoxy-4-dedimethylaminotetracycline, 12a, 4  $\alpha$  -anhydro-4-dedimethylaminotetracycline, 7-dimethylamino-6-demethyl-6-deoxy-4-dedimethylaminotetracycline, 5-hydroxy-6-  $\alpha$  -deoxy-4-dedimethylaminotetracycline, 4-dedimethylamino-12  $\alpha$  -deoxyanhydrotetracycline and 4-dedimethylamino-11-hydroxy-12a-deoxytetracycline.

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B<sup>2</sup> 27. (Amended) A method according to Claim 18, wherein said tetracycline derivative is doxycycline.

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28. A method according to Claim 18, wherein said tetracycline derivative is administered systemically.

29. A method according to Claim 28, wherein said tetracycline derivative is administered systemically by a controlled release delivery system.